Product Data Sheet

isotype control (open histogram).

Brilliant Violet 605[™] anti-human HLA-DR

Catalog # / Size:	2138200 / 100 tests 2138195 / 25 tests	A
Clone:	L243	18
Isotype:	Mouse IgG2a, к	la
Reactivity:	Human	N N
Preparation:	The antibody was purified by affinity chromatography and conjugated with Brilliant Violet 605 [™] under optimal conditions. The solution is free of unconjugated Brilliant Violet 605 [™] and unconjugated antibody.	Relative Cell Number
Formulation:	Phosphate-buffered solution, pH 7.2, containing 0.09% sodium azide and BSA (origin USA).	Log Fluorescence Intensity Human peripheral blood lymphocytes were stained with HLA- DR (clone L243) Brilliant Violet 605 [™] (filled histogram) or mouse lgG2a, κ Brilliant Violet 605 [™]
Concentration:	Lot-specific	

Applications:

Applications: Flow Cytometry	Appl	ications:	Flow Cytometr	y
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Each lot of this antibody is quality control tested by immunofluorescent staining Recommended with flow cytometric analysis. For flow cytometric staining, the suggested use of Usage: this reagent is \leq 5 microL per million cells or 5 microL per 100 microL of whole blood. It is recommended that the reagent be titrated for optimal performance for each application.

> Brilliant Violet 605[™] excites at 405 nm and emits at 603 nm. The bandpass filter 610/20 nm is recommended for detection, although filter optimization may be required depending on other fluorophores used. Be sure to verify that your cytometer configuration and software setup are appropriate for detecting this channel. Refer to your instrument manual or manufacturer for support. Brilliant Violet 605[™] is a trademark of Sirigen Group Ltd.

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Application The L243 monoclonal antibody reacts with the HLA-DR antigen, a member of MHC class II molecules. It does not cross react with HLA-DP and HLA-DQ. Clone L243 Notes: binds a conformational epitope on HLA-DR α which depends on the correct folding of the $\alpha\beta$ heterodimer.¹⁹

Additional reported applications (for the relevant formats) include: immunoprecipitation⁸, Western blotting⁸, *in vitro* blocking of mixed lymphocyte reactions^{9,10}, depeletion of MHC class II cells⁷, and immunohistochemical staining of acetone-fixed frozen sections^{4,5}. The LEAF $^{\text{\tiny M}}$ purified antibody (Endotoxin <0.1 EU/µg, Azide-Free, 0.2 µm filtered) is recommended for functional assays (Cat.

No. 307612). For highly sensitive assays, we recommend Ultra-LEAF^m purified antibody (Cat. No. 307648) with a lower endotoxin limit than standard LEAF^m purified antibodies (Endotoxin <0.01 EU/microg).

Application References:	 Brodsky F. 1984. <i>Immunogenetics</i> 19:179. Robbins P, <i>et al.</i> 1987. <i>Human Immunol.</i> 18:301. Stites D, <i>et al.</i> 1986. <i>Clin. Immunol. Immunopathol.</i> 38:161. Warnke R, <i>et al.</i> 1980. <i>J. Histochem. Cytochem.</i> 28:771. (IHC) Engleman E, <i>et al.</i> 1981. <i>P. Natl. Acad. Sci. USA</i> 78:1791. (IHC) Zipf T, <i>et al.</i> 1981. <i>Cancer Res.</i> 41:4786. Goodier M, <i>et al.</i> 2000. <i>J. Immunol.</i> 165:139. (Depletion) Esser M, <i>et al.</i> 2001. <i>J. Virol.</i> 75:6173. (IP, WB) Kalka-Moll WM, <i>et al.</i> 2002. <i>J. Immunol.</i> 169:6149. (Block) Wang RF, <i>et al.</i> 1999. <i>Science</i> 284:1351. (Block) Vang RF, <i>et al.</i> 2007. <i>J. Exp. Med.</i> 204:3183. PubMed Fujita H, <i>et al.</i> 2000. <i>P. Natl. Acad. Sci. USA</i> 106:21795. PubMed Charles N, <i>et al.</i> 2010. <i>Infect. Immun.</i> 78:4763. PubMed Yoshino N, <i>et al.</i> 2000. <i>Exp. Anim. (Tokyo)</i> 49:97. (FC) Kim WK, <i>et al.</i> 2001. <i>Leuk. Lymphoma</i> 52:273. Galkowska H, <i>et al.</i> 1996. <i>Vet. Immunol.</i> 169:819. PubMed Salkowska H, <i>et al.</i> 2005. <i>BMC Immunol.</i> 59:19. PubMed
Description:	HLA-DR is a heterodimeric cell surface glycoprotein comprised of a 36 kD $lpha$

Description:	HLA-DR is a heterodimeric cell surface glycoprotein comprised of a 36 kD $lpha$
	(heavy) chain and a 27 kD β (light) chain. It is expressed on B cells, activated T
	cells, monocytes/macrophages, dendritic cells, and other non-professional APCs.
	In conjunction with the CD3/TCR complex and CD4 molecules, HLA-DR is critical
	for efficient peptide presentation to CD4 ⁺ T cells.

Antigen	1. Levacher M, <i>et al.</i> 1990. <i>Clin. Exp. Immunol.</i> 81:177.	
References:	2. Terstappen L, et al. 1990. J. Leukocyte Biol. 48:138.	
	3. Edwards JA, <i>et al.</i> 1986. <i>J. Immunol.</i> 137:490.	
	4. van Es A, <i>e</i>	

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